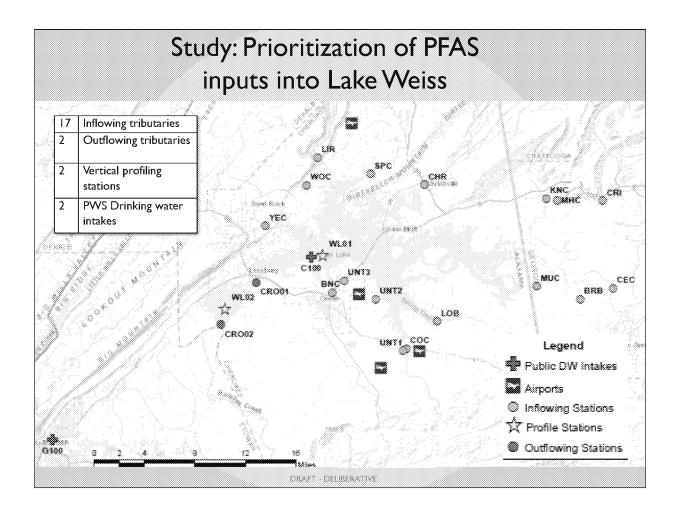
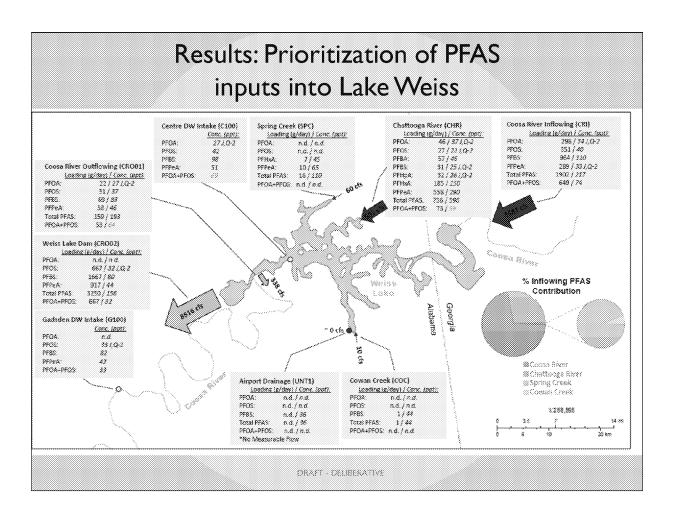


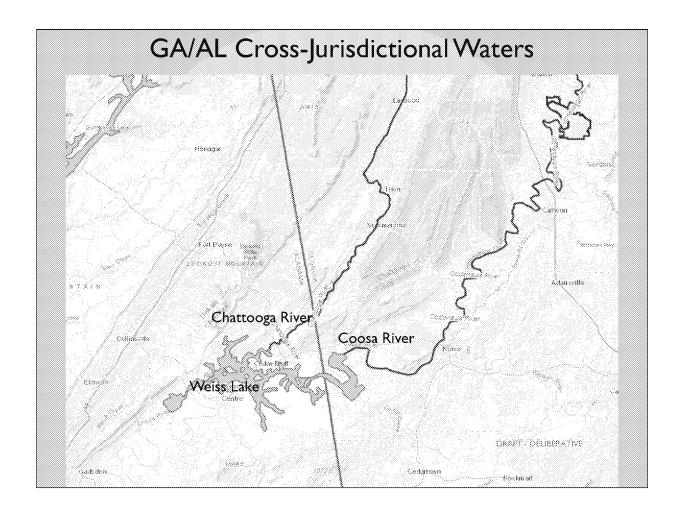
PRIORITIZATION OF PFAS CONTRIBUTIONS TO WEISS LAKE

Study Objectives:

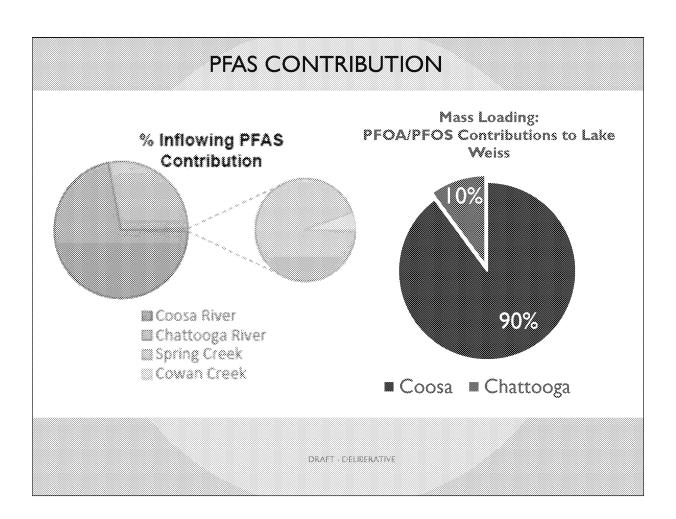
- Measure background PFAS concentrations and streamflow measurements for all inflowing and outflowing tributary station of Weiss Lake.
- 2. Estimate mass loading rates of PFASs into Weiss Lake.
- 3. Examine vertical distribution of PFASs within the lake water column and understand the correlation with water quality parameters.

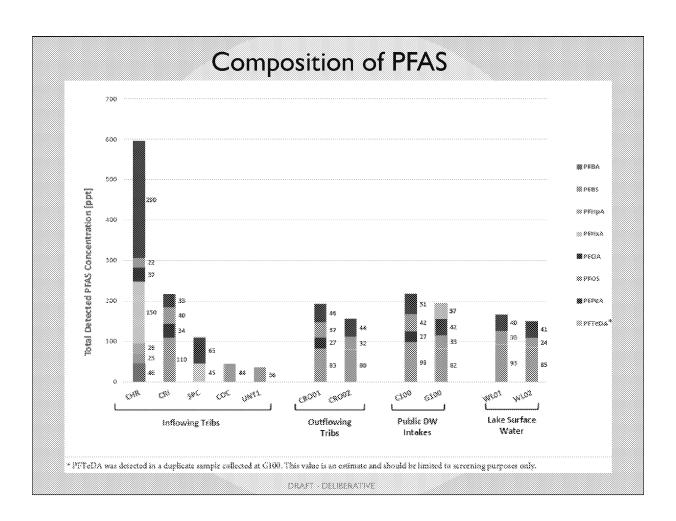


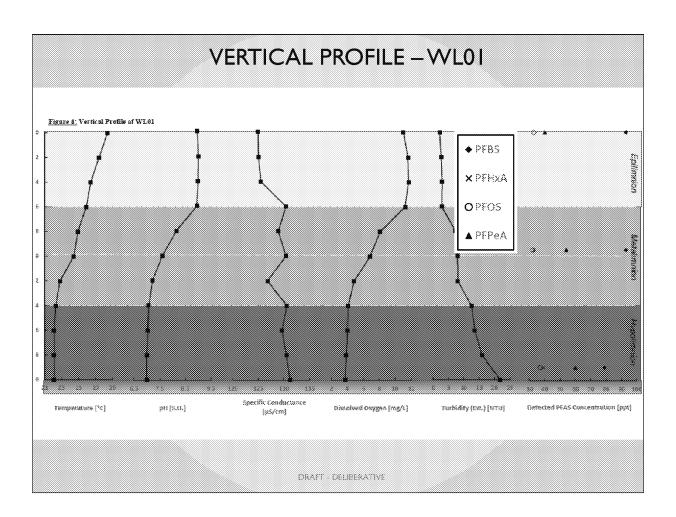


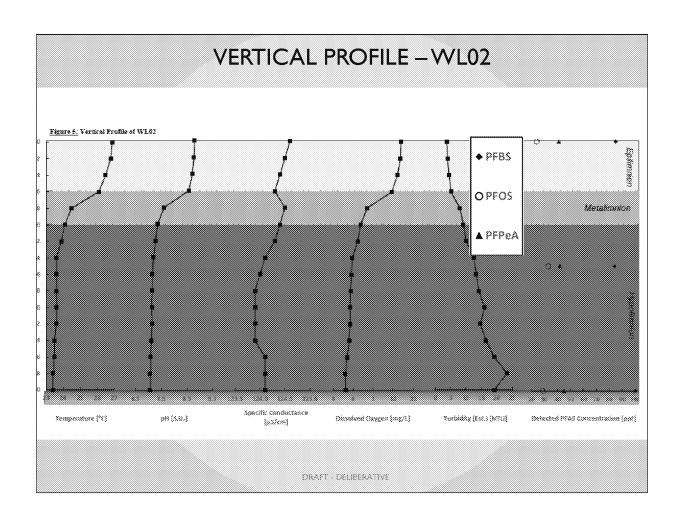


13 co-located SW and sediment samples from LaFayette through Weiss Lake









PRIORITIZATION OF PFAS CONTRIBUTIONS TO WEISS LAKE

Conclusions:

- Coosa River inlet sample
 - 74 ppt (combined PFOA/PFOS)
 - Largest mass loading rate 7x flow from Chattooga
 - 90% of combined PFOA/PFOS; 72% of total PFAS
- Chattooga River inlet sample
 - 59 ppt (combined PFOA/PFOS)
 - Greater diversity of PFASs and largest total PFAS concentration
- Consistent distribution of 3 PFASs (PFOS, PFPeA, PFBS) vertically (up to 30 feet below surface) throughout a thermally stratified water column.
 - PFAS contributions from surface water and sediment samples
- Most prevalent PFASs: PFPeA and PFBS (short-chain compounds)

Question: Can we use the LHA as reference to surface water PFOA/PFOS?



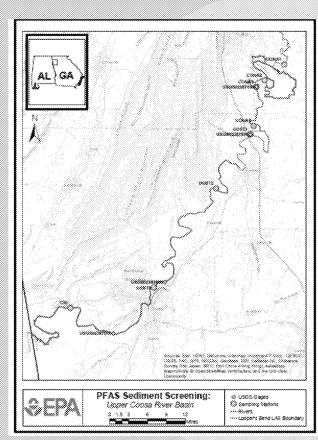
ASSESSMENT OF RESUSPENDED SEDIMENTS AS A SOURCE OF PFAS TO THE UPPER COOSA RIVER BASIN

Conasauga, Oostanaula and Coosa Rivers

Study Objectives:

- 1. Measure PFAS concentration and composition in stream sediments impacted by the Looper's Bend LAS along the Conasauga into the receiving waters of the Oostanaula and Coosa Rivers.
- 2. Compare PFAS concentration and composition in co-located surface water and sediment samples.
- 3. Estimate instantaneous suspended sediment flux to determine the downstream PFAS migration of sediments.

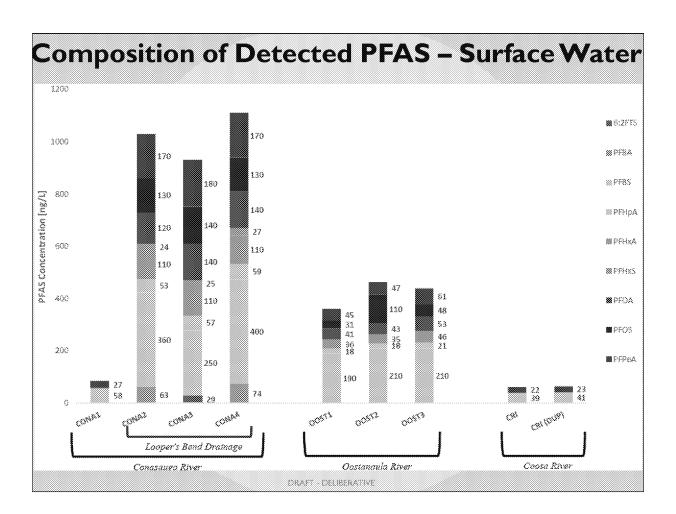
^{*}Targeted base-flow conditions to avoid PFAS Inputs from surface water runoff and dilution

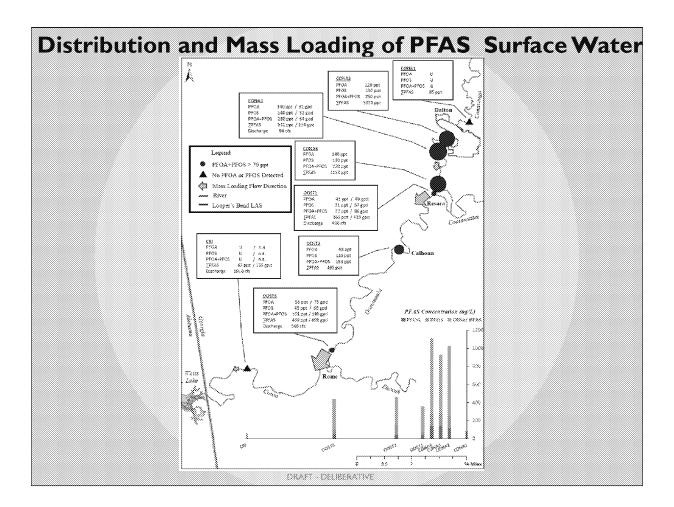


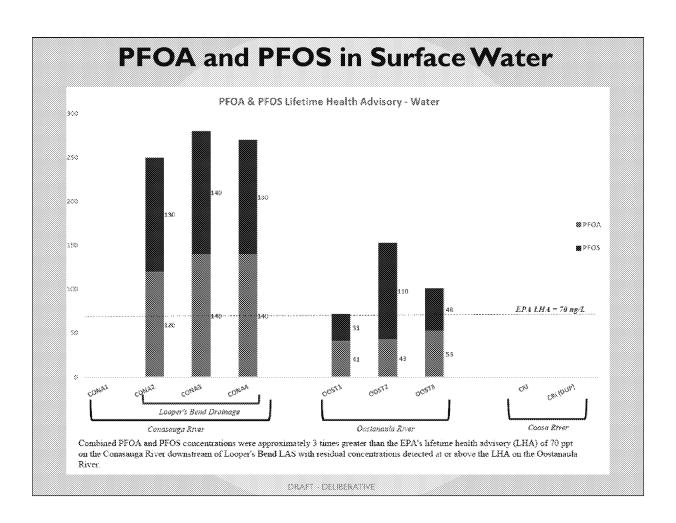
Total of 8 sites sampled for 25 PFAS Analytes:

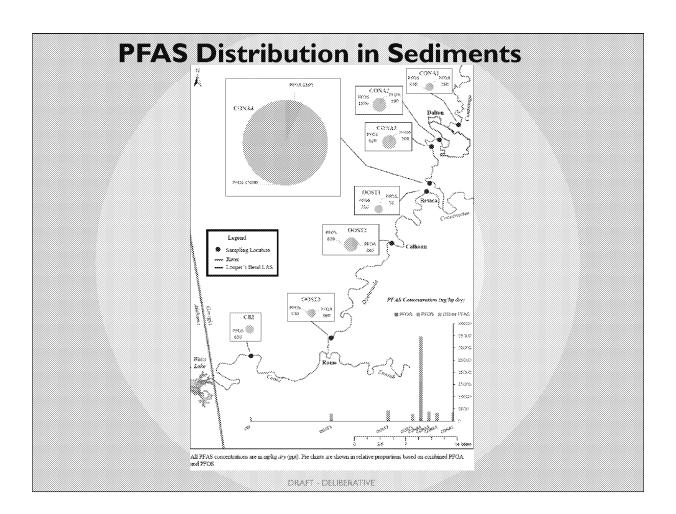
- 8 sites co-located sediments and surface waters
 - TOC & TSS (sediment)
- 4 of the 8 surface water samples included the following additional analysis:
 - Filtration to determine fraction of PFAS suspended in water column
 - USGS Streamflow
 - TOC & TSS (surface water)

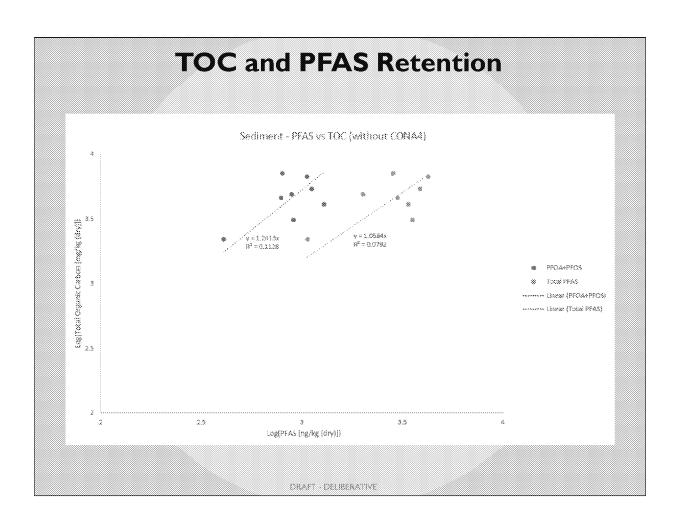
DRAFT CELEERATIVE







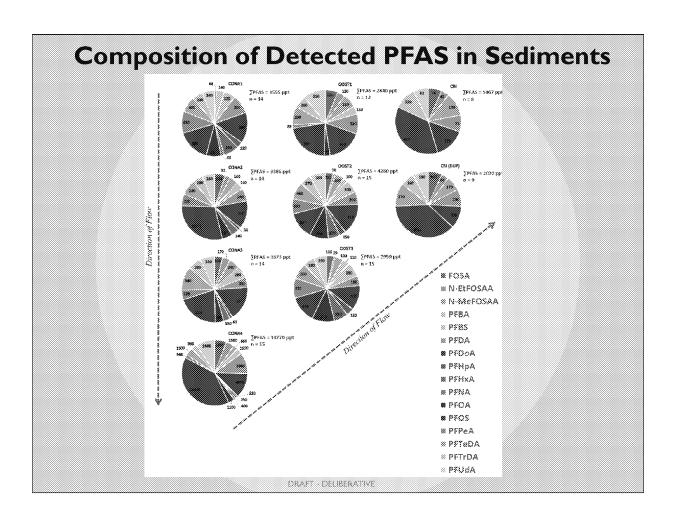


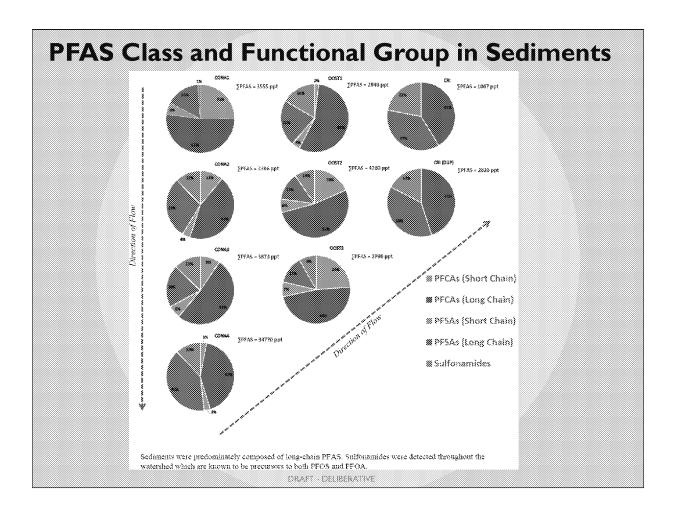


Functional Group and Classification

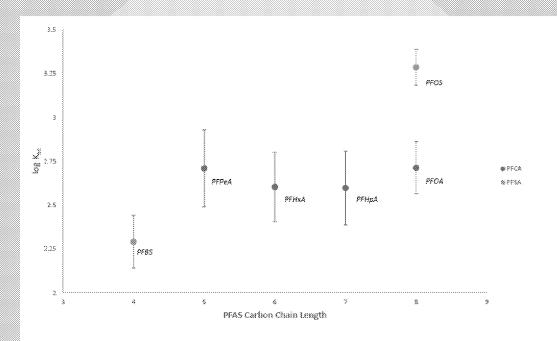
Carbon Chain Length	PFAS Analyte	Class	Functional Group
14	PFTeDA	Long Chain	PFCAs
13	PFTrDA		
12	PFDoA		
11	PFUDA		
10	PFDA		
9	PFNA		
8	PFOA		
-	PEHpA	Short Chain	
6	PFHA		
•	PIPeA		
4	PFBA		
10	PFDS	Long Chain	PFSAs
ŷ.	PFNS		
8	PFOS		
7	PFHpS		
б	PFHxS		
	PFPeS	Short Chain	
4	PEBS		
10	8:2 FTS	Precursor	Fluorotelomers
8	6:2 FTS		
- 6	4:2 FTS		
10	N-EIFOSAA	Precursor	Sulfonamides
9	N-MeFOSAA		
- 8	FOSA		
	DEPOSITS OF	0.000	Distriction

Classifications adapted from ITRC, 2018



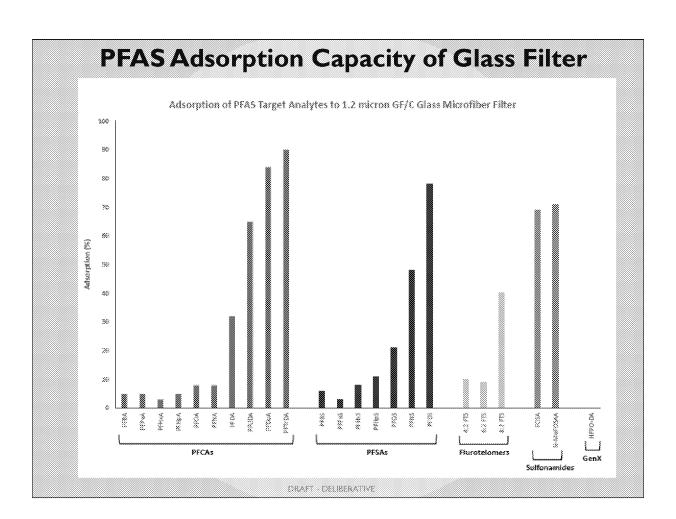


Organic Carbon on PFAS Retention in Sediments



The effect of organic carbon and the retention of PFAS in sediments is dependent on both the chain length and functional group. PFAS retention in sediments generally increases with increasing chain length and PFSA3 (e.g. PFOS) are more likely to be retained compared to PFCAs (e.g. PFOA) of the same chain length.

CRAFT CELESRATIVE



UPPER COOSA RIVER BASIN: SURFACE WATER-SEDIMENTS

Conasauga, Oostanaula and Coosa Rivers

Conclusions:

Surface water:

- PFBS and PFPeA (short-chain compounds) are most prevalent throughout the watershed (similar to Prioritization Study)
- Generally dominated by short-chain PFSAs and PFCAs (highly water-soluble)

Sediment:

- Dominated by long-chain PFCAs, followed by long-chain PFSAs and sulfonamido precursors (higher adsorption capacity)
- Total PFAS influenced by the presence of TOC

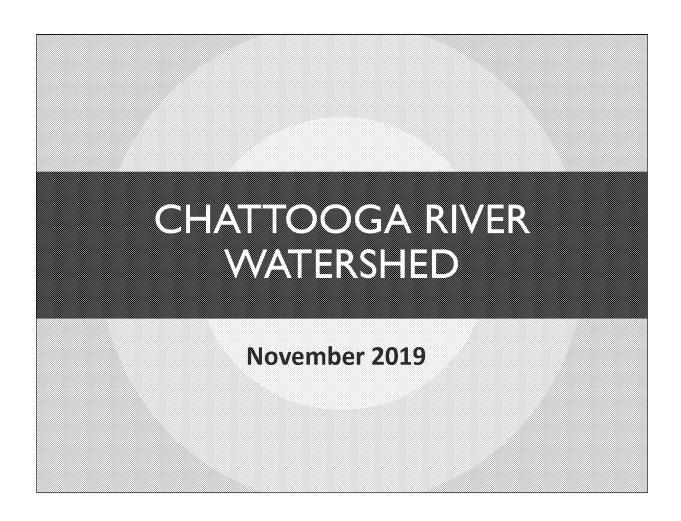
UPPER COOSA RIVER BASIN: SURFACE WATER-SEDIMENTS

Conasauga, Oostanaula and Coosa Rivers

Conclusions:

Mass Loading:

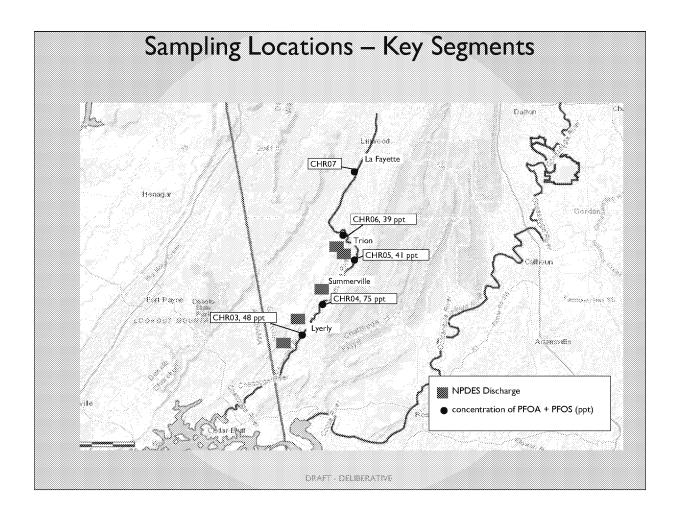
- Conasauga River: 74% of PFOA/PFOS Mass Loading to the Oostanaula River
- Mass loading analysis limited by current Method Detection Limit
- Highest diversity and concentrations of PFAS observed downstream of Looper's Bend persisting in the receiving waters
- The contribution of resuspended sediments was inconclusive due to high variability and adsorption loss of PFAS to filters



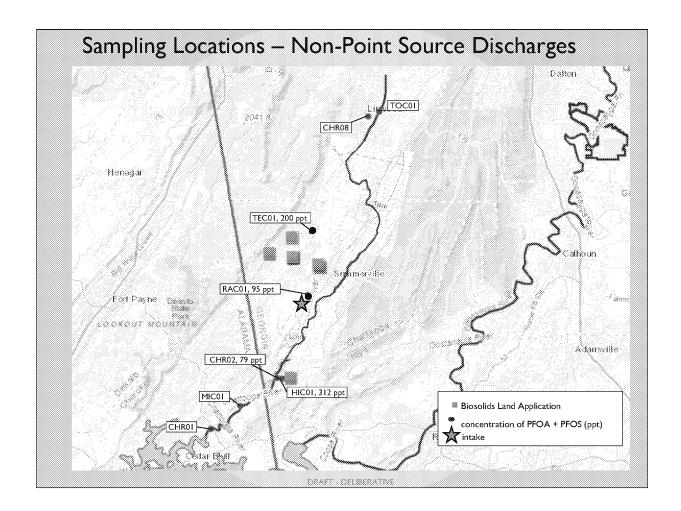
Characterization of Ambient PFAS in the Chattooga River Watershed

Study Objectives:

- Characterize the distribution and instantaneous mass loading of PFAS in the Chattooga River Watershed at near-base flow conditions along key segments determined by the R4 water division.
- 2. Collect surface water samples coupled with discharge measurements to compute instantaneous mass loading rates of PFAS along key segments of the Chattooga River Watershed.
- 3. Collect sediment samples collocated with surface water sample locations to determine the relative distribution and the potential for migration of PFAS contaminated sediments to the receiving waters of

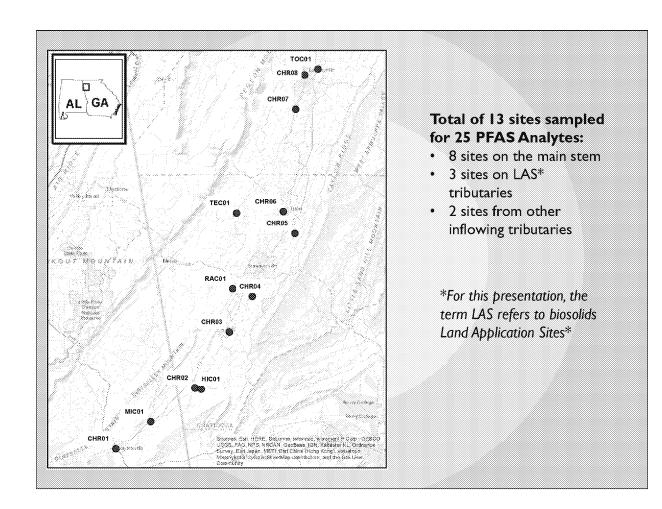


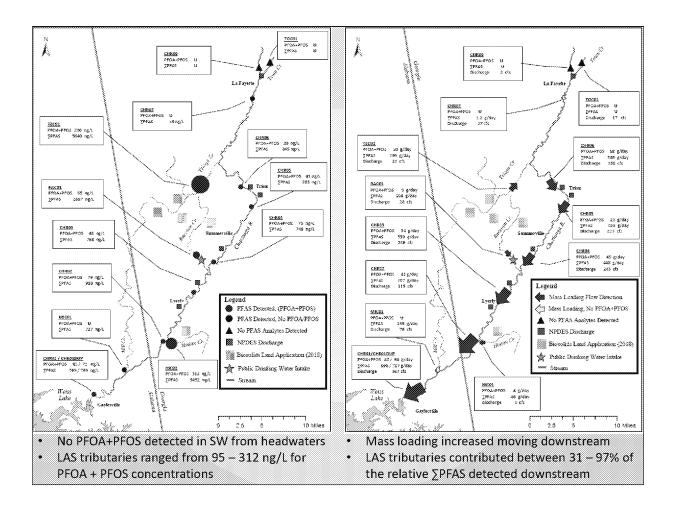
6 SW samples had results above the LHA

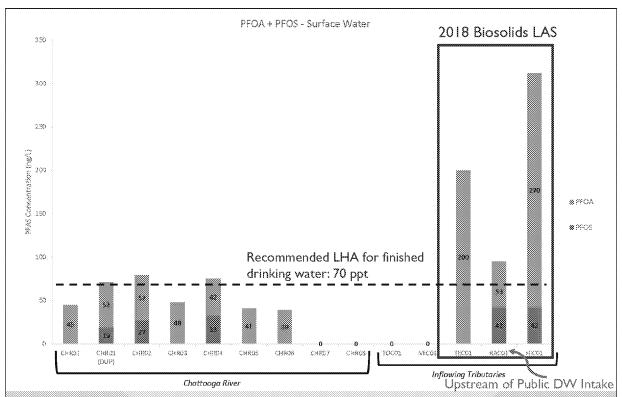


Intakes, biosolids fields and SW samples above LHA

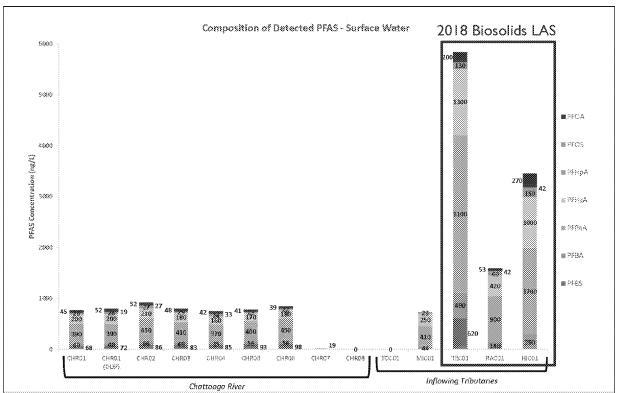
Unlike the Coosa River watershed, more biosolids fields are in the Chattooga



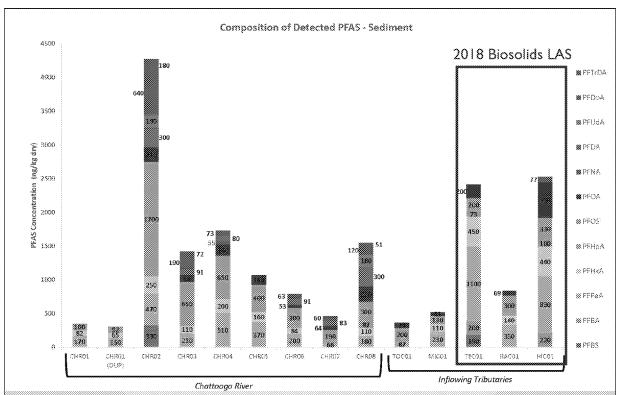




- Surface water sample results represent a snapshot in time
- TEC01, RAC01, & HIC01 are tributary sites impacted by biosolids land application
- Public drinking water intake for City of Summerville located just upstream of station RAC01
- Surface waters dominated by short-chain PFCAs



- Surface water sample results represent a snapshot in time
- TEC01, RAC01, & HIC01 are tributary sites impacted by biosolids land application
- Public drinking water intake for city of Summerville located just upstream of station RAC01
- Surface waters dominated by short-chain PFCAs



- Sediment sample results represent integration over time & are dependent on site conditions, such as the amount of total organic carbon (TOC) present
- TEC01, RAC01, & HIC01 are tributary sites impacted by biosolids land application
- Sediments generally dominated by short-chain PFCAs

Characterization of Ambient PFAS in the Chattooga River Watershed

Conclusions

Sediment Sampling Results: 12 distinct PFAS compounds detected, <u>majority</u> of samples dominated by short-chain PFCAs

<u>Surface Water Sampling Results:</u> 7 distinct PFAS compounds detected, <u>all</u> samples dominated by short-chain PFCAs

- · No PFAS detected in surface water samples from headwaters
- Significant contribution of ∑PFAS & combined PFOA+PFOS inputs from tributaries with active biosolids land application sites
- Short-chain PFCAs (C5&C6) dominate both SW & SD, suggesting a local source of short-chain PFCAs &/or their precursors within the watershed
- * Transport of both SW & SD contaminated with PFAS from the Chattooga River Watershed \rightarrow Weiss Lake DRAFT DELBERATIVE